A method of inhibiting mammalian hair growth 1 2 which comprises

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selecting an area of skin from which reduced hair growth is desired; and

applying to said area of skin a dermatologically acceptable composition comprising a non-steroidal suppressor of angiogenesis in an amount effective to inhibit hair growth.

- The method of claim 1, wherein said suppressor 1 is a compound that interferes with the action of heparin 2 3 sulfate proteoglycans.
- The method of claim 2, wherein said compound is 1 an inhibitor of sulfotransferase. 2
- The method of claim 2, wherein said compound is 1 a heparin binding antagonist. 2
- The method of claim 2, wherein said compound is 1 2 a copper chelator.
- The method of claim 1, wherein said suppressor 1 is a compound that interferes with the action of histamine. 2
- The method of claim 6, wherein said compound is 1 an inhibitor of histidine decarboxylase.
  - The method of claim 6, wherein said compound is an inhibitor of mast cell degranulation.

- 9. The method of claim 6, wherein said compound is a histamine receptor antagonist.
- 1 10. The method of claim 1, wherein said suppressor
- 2 is a compound that interferes with the action of angiotensin
- 3 II.
- 1 11. The method of claim 10, wherein said compound
- 2 is an inhibitor of angiotensin converting enzyme.
- 1 12. The method of claim 10, wherein said compound
- 2 is an angiotensin II receptor antagonist.
- 1 13. The method of claim 1, wherein said suppressor
- 2 is a compound that interferes with the action of
- 3 prostaglandin E1.
  - 14. The method of <u>claim 1</u>, wherein said compound is an inhibitor of prostaglandin synthetase.
- 1 15. The method of claim 1 wherein said suppressor
- 2 interferes with the action of Substance P.
- 1 16. The method of claim 15, wherein said compound
- 2 is an NK1 receptor antagonist.
- 1 17. The method of claim 1, wherein said suppressor
- 2 interferes with the action of platelet activating factor.
- 1 18. The method of claim 17, wherein said compound
- 2 is a platelet activating factor receptor antagonist.

- 1 19. The method of claim 1, wherein said suppressor
- 2 interferes with the action of 12-HETrE.
- 1 20. The method of claim 19, wherein said compound
- 2 is an inhibitor of cytochrome P450 reductase.
- 1 21. The method of claim 1, wherein said composition
- 2 further comprises vehicle.
- 1 22. The method of claim 1, wherein the
- 2 concentration of said suppressor in said composition is
- 3 between 1% and 30% by weight.
- 1 23. The method of claim 1, wherein the composition
- 2 provides a reduction in hair growth of at least 20% when
- 3 tested in the Golden Syrian hamster assay.
- 1 24. The method of claim 1, wherein the composition
- 2 provides a reduction in hair growth of at least 50% when
- 3 tested in the Golden Syrian hamster assay.
- 1 25. The method of claim 1, wherein the composition
- 2 provides a reduction in hair growth of at least 70% when
- 3 tested in the Golden Syrian hamster assay.
- 1 26. The method of claim 1, wherein the suppressor
- 2 is applied to the skin in an amount of from 100 to 3000
- 3 micrograms of said inhibitor per square centimeter of skin.
- 1 27. The method of claim 1, wherein said mammal is a human.

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1	The method of claim 27, wherein said area of
2	skin is on the face of the human.
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1	29. The method of claim 28, wherein said human is a
2	woman suffering from hirsutism.
1	reducing 30. A method of inhibiting mammalian hair growth
2	which comprises
_3	selecting an area of skin from which reduced hair
$\geq$ 4	growth is desired; and $ackslash$
5	applying to said area of skin a dermatologically
6	acceptable composition comprising an inhibitor of
7	sulfotransferase in an amount effective to inhibit hair
8	growth.
	20 29
1	The method of claim 30, wherein said inhibitor
2	is p-nitrocatechol.
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1	37. The method of <u>claim 30</u> , wherein said inhibitor
2	is catechin.
	reducin a
1	reducing  33. A method of inhibiting mammalian hair growth
2	which comprises
3	selecting an area of skin from which reduced hair
>4	growth is desired; and $igwedge$
5	applying to said area of skin a dermatologically
6	acceptable composition comprising a heparin binding
7	antagonist in an amount effective to inhibit hair growth.
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1	37. The method of claim 33, wherein said antagonist
2	is pentosan polysulfate.

The method of claim 23, wherein said antagonist 1 2 is quinacrine. A method of inhibiting mammalian hair growth 1 2 which comprises selecting an area of skin from which reduced hair 3 growth is desired; and applying to said area of skin a dermatologically acceptable composition comprising a copper chelator in an 6 amount effective to inhibit hair growth. 7 The method of claim 36, wherein said copper 1 2 chelator is bathocuproine disulfonate. The method of claim 36, wherein said copper 1 2 chelator is diethylenetriamine pentaacetic acid. A method of inhibiting mammalian hair growth 1 2 which comprises 3 selecting an area of skin from which reduced hair growth is desired; and applying to said area of skin \a dermatologically acceptable composition comprising an inhibitor of histidine 6 decarboxylase in an amount effective to inhibit hair growth. 7 The method of claim 39, wherein said inhibitor 1 2 is O-p-nitrohydroxylamine.

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is  $\alpha$ -fluoromethylhistidine.

The method of claim 39, wherein said inhibitor

reducina A method of inhibiting mammalian hair growth 1 42. 2 which comprises selecting an area of skin from which reduced hair 3 growth is desired; \and applying to said area of skin a dermatologically 5 acceptable composition comprising an inhibitor of mast cell 6 degranulation in an amount effective to inhibit hair growth. 7 The method of claim 42, wherein said inhibitor 1 2 is mycophenolic acid. The method of claim 42, wherein said inhibitor 1 2 is bromocryptine. 1 The method of claim A2, wherein said inhibitor 2 is cromoglycate. A method of inhibiting mammalian hair growth 1 2 which comprises selecting an area of skin from which reduced hair growth is desired; and applying to said area of skin a dermatologically 6 acceptable composition comprising a histamine receptor antagonist in an amount effective to inhibit hair growth. 7 1 The method of claim 46, wherein said antagonist 2 is terfenadine.

The method of claim 46, wherein said antagonist

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is tripélennamine.

The method of claim 46, wherein said antagonist 1 2 is chlorpheniramine. The method of claim A6, wherein said antagonist 1 is cimetidine. 2 A method of inhibiting mammalian hair growth 1 which comprises 2 selecting an area of skin from which reduced hair growth is desired; and 5 applying to said area of skin a dermatologically acceptable composition comprising an inhibitor of 6 angiotensin converting enzyme in an amount effective to inhibit hair growth. 8 The method of claim 51, wherein said inhibitor 1 2 is enalapril. The method of claim 51, wherein said inhibitor 1 2 is lisinopril. A method of inhibiting mammalian hair growth 1 2 which comprises selecting an area\of skin from which reduced hair growth is desired; and 5 applying to said area of skin a dermatologically acceptable composition comprising an angiotensin II receptor 6 antagonist in an amount effective to inhibit hair growth. 7 The method of claim 34, wherein said antagonist 1

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is a 1,4- substituted indole.

The method of claim 54, wherein said antagonist 1 is a dihydropyridine derivative. 2 The method of claim 56, wherein said antagonist 1 is nifedipine. 2 The method of claim 54, wherein said antagonist 1 is a triazolinone derivative. 2 The method of claim 58, wherein said 1 triazolinane derivative has a side chain at the N4 position. 2 The method of claim 54, wherein said antagonist 3 6Ó. is a tetrahydroisoquinoline carboxylic acid. 4 60 The method of claim 54, wherein said antagonist 5 6 is an imidazopyridine derivative. 101 The method of claim 61, wherein said 7 imidazopyridine derivative is a tetrahydroimidazopyridine 8 carboxylic acid analog. 9 The method of claim 54, wherein said antagonist 10

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is Losantan.

reducina A method of inhibiting mammalian hair growth 1 2 which comprises 3 selecting an area of skin from which reduced hair growth is desired; and applying to said area of skin a dermatologically acceptable composition comprising an inhibitor of 7 prostaglandin synthetase in an amount effective to inhibit 8 hair growth. 65. The method of claim 64, wherein said inhibitor is piracetam. reducing A method of inhibiting mammalian hair growth 1 2 which comprises selecting an area of skin from which reduced hair 3 growth is desired; and applying to said area of skin a dermatologically acceptable composition comprising an NK1 receptor antagonist in an amount effective to inhibit hair growth. 7 1 The method of claim 6%, wherein said antagonist 2 is (3aR,7aR)-7,7,-diphenyl-2-[1-imino-2-(2-methoxyphenyl) ethyl]perhydroisoindol-4-one. 3 The method of claim 6%, wherein said antagonist 1 is cis-2-(diphenylmethyl)-N-[(2-methoxy-phenyl)]-methyl]-1-2

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azabicyclo[2,2,2]octan-3-amine.

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A method of inhibiting mammalian hair growth 1 2 which comprises selecting an area of skin from which reduced hair 3 growth is desired; and applying to said area of skin a dermatologically acceptable composition comprising a platelet activating factor receptor antagonist in an amount effective to inhibit 7 hair growth. 8 The method of claim 6%, wherein said antagonist 1 2 is tioconazole. 10 71. The method of claim 69, wherein said antagonist 1 is (3-[4-(2-chlorphenyl)-9-methyl-6H-thieno[2-f]-2 [1,2,4]triazolo-[4,3-a][1,4]-diazepin-2-yl-1-(4-3 morpholinyl)-1-propanone. A method of inhibiting mammalian hair growth 1 2 which comprises selecting an area of skin from which reduced hair growth is desired; and applying to said area of skin a dermatologically acceptable composition comprising an inhibitor of cytochrome 6 P450 reductase in an amount effective to inhibit hair 7 growth. The method of claim 22, wherein said inhibitor 1 is clotrimazole.

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74. A method of inhibiting mammalian hair growth which comprises

selecting an area of skin from which reduced hair growth is desired; and

applying to said area of skin a dermatologically acceptable composition comprising, in an amount effective to reduce hair growth, a compound selected from the group consisting of phenyl-ethylene derivatives such as tamoxifen and nafoxidine; irsogladine; the synthetic laminin peptide,

10 CDPGYIGSR-NH2; radicicol; eponemycin; fumagillin (0-

11 (chloroacetyl-carbamoyl) fumagillol) and synthetic analoges

12 thereof; recombinant\human platelet factor-4 and related

13 peptides; protamine; sulfated chitin derivatives;

14 diaminoanthraquinone derivatives; thrombospondin; quinoline-

15 3-carboxamide (linomide); analogues of diatamycin A; and

16 aurintricarboxylic acid.

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